If you have IRON DEFICIENCY ANEMIA (IDA) and have an intolerance or an unsatisfactory response to oral iron,

INJECTAFER WORKS HARD TO REPLENISH YOUR IRON LEVELS

See what Injectafer can do

Injectafer® (ferric carboxymaltose injection) is available by prescription only. Ask your healthcare provider if Injectafer is right for you.

What is Injectafer?

Injectafer is a prescription iron replacement medicine administered only by or under the supervision of your healthcare provider. Injectafer is injected into your vein to treat iron deficiency anemia in adults and children 1 year of age and older. Injectafer should be used only if you have not responded well to treatment with oral iron, or if you are intolerant to oral iron treatment. It is also used to treat iron deficiency anemia in adults with chronic kidney disease who are not receiving dialysis. Injectafer is used to improve the ability to exercise (exercise capacity) in adult patients with iron deficiency and mild to moderate heart failure. It is not known if Injectafer is safe and effective in children with iron deficiency anemia who are under 1 year of age or in children with iron deficiency and mild to moderate heart failure to improve exercise capacity.

Please click here for Full Prescribing Information and see Important Safety Information on pages 18-19.

injectafer®
ferric carboxymaltose injection

GET TO KNOW INJECTAFER

AN INTRAVENOUS (IV) IRON TREATMENT FOR IRON DEFICIENCY ANEMIA (IDA)

Ask your healthcare professional to check your iron levels!

SELECTED SAFETY INFORMATION

Who should not receive Injectafer?

You should not receive Injectafer if you are allergic to ferric carboxymaltose or any of the other ingredients in Injectafer.

The active ingredient in Injectafer is ferric carboxymaltose, the inactive ingredients are: water for injection, sodium hydroxide or hydrochloric acid.

CLICK HERE TO VISIT INJECTAFER.COM/INFO

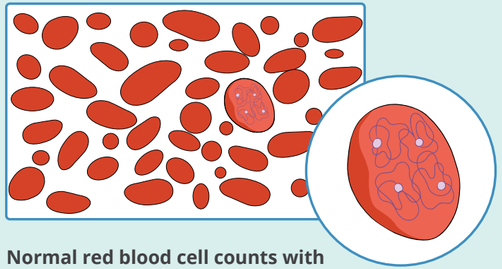
*Terms and conditions apply.

Please click here for Full Prescribing Information and see Important Safety Information on pages 18-19.

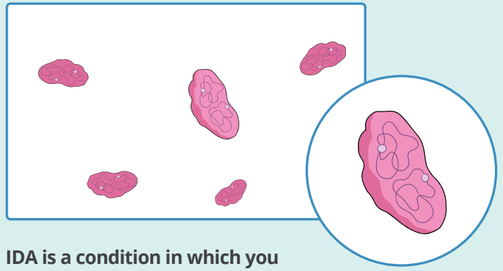
injectafer®
ferric carboxymaltose injection

WHAT IS IDA?

IRON DEFICIENCY ANEMIA (IDA) IS A CONDITION THAT INTERFERES WITH THE FORMATION AND FUNCTION OF RED BLOOD CELLS



Normal red blood cell counts with a healthy level of oxygen-carrying hemoglobin (Hb)



IDA is a condition in which you have fewer healthy red blood cells and less Hb inside each cell

SELECTED SAFETY INFORMATION

What should I tell my healthcare provider before receiving Injectafer® (ferric carboxymaltose)?

Before you receive Injectafer, tell your healthcare provider about all of your medical conditions, including if you:

- Have had an allergic reaction to iron given into your vein
- Have a history of trouble absorbing certain vitamins or phosphate in your body
- Have inflammatory bowel disease

WHAT CAUSES IDA?

1

Iron is either depleted or not fully replenished due to causes such as blood loss, lack of dietary iron, inability to absorb iron, or pregnancy.

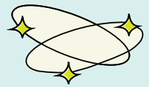
2

Low iron levels (also called iron deficiency) occur when you don't have enough iron to keep you in good health.

3

If iron deficiency progresses, it may cause IDA, as iron helps produce hemoglobin (Hb).

WHAT ARE SOME OF THE SYMPTOMS OF IDA?*



DIZZINESS

WEAKNESS

PICA
(craving non-food items such as ice or dirt)

FATIGUE

SHORTNESS OF BREATH

Some people with IDA experience a range of symptoms, while other people experience no symptoms at all. Not all patients feel the same way with IDA. It's possible to confuse the signs of iron deficiency anemia with symptoms of other conditions you may have.

*Injectafer has not been studied or approved for treating symptoms of IDA.

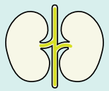
Please click here for Full Prescribing Information and see Important Safety Information on pages 18-19.

MANY DISEASES AND CONDITIONS MAY CAUSE OR LEAD TO IDA

WOMEN'S HEALTH CONDITIONS

GASTROINTESTINAL CONDITIONS

CANCER



CHRONIC KIDNEY DISEASE

HEART FAILURE

BLOOD LOSS FROM INJURY OR SURGERY

HOW IS IDA DIAGNOSED?

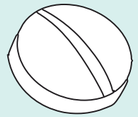
IDA is determined by testing 3 different lab markers in your blood: **hemoglobin (Hb), ferritin, and transferrin saturation (TSAT).***

At your next appointment, ask your healthcare professional when the last time your iron levels (including Hb, ferritin, and TSAT) were checked. **If it was more than 3 months ago, ask them to check your levels again.**

*An additional test for total iron binding capacity (TIBC) values may be required to diagnose IDA.

Please click here for Full Prescribing Information and see Important Safety Information on pages 18-19.

HOW IS IDA TREATED?



ORAL IRON

OR

IV IRON

When treating IDA, both oral and IV iron are common treatment options. There are several reasons why oral iron supplements may not be suitable for some people with IDA:

- Iron supplements can cause **hard-to-tolerate side effects**
- **Poor absorption:** the digestive tract is only able to absorb a small portion of the iron in an iron supplement, so your body may not get the full dose of iron needed*

SPEAK UP AT YOUR NEXT APPOINTMENT

If you are experiencing any symptoms while taking oral iron, no matter how mild they may be, tell your healthcare professional.

*Oral iron is typically taken in 300 mg or 320 mg tablets 3-4 times a day. The body is unable to absorb that much iron in the digestive tract at one time, so iron repletion may be possible in smaller oral iron doses over time.

SELECTED SAFETY INFORMATION

What should I tell my healthcare provider before receiving Injectafer® (ferric carboxymaltose)? (cont'd)

Before you receive Injectafer, tell your healthcare provider about all of your medical conditions, including if you:

- Have hyperparathyroidism
- Have low vitamin D levels
- Have high blood pressure
- Have previously received Injectafer

UNDERSTANDING YOUR LAB VALUES

After your healthcare professional orders a blood test, you will receive the lab results and you'll likely want to know more about what they mean.

This chart can help you understand your lab results. The chart shows "normal" values but keep in mind that these can vary for many reasons, depending on the person, medical condition(s), and where the test was taken.

Lab markers	Normal values in healthy adults	Normal values in healthy children
HEMOGLOBIN (Hb)	13.5-17.5 g/dL (for males) 12.0-15.5 g/dL (for females)	10.5-16 g/dL (for all children)
FERRITIN	40-300 ng/mL (for males) 20-200 ng/mL (for females)	36-311 ng/mL (for males) 36-92 ng/mL (for females)
TRANSFERRIN SATURATION (TSAT)	20%-50% (for all adults)	15%-44% (for males) 11%-44% (for females)

GET YOUR LEVELS CHECKED

If your lab values are below normal range, talk to your healthcare professional about the best option for restoring your iron levels.

Please click here for Full Prescribing Information and see Important Safety Information on pages 18-19.

HOW DO I KNOW IF I'M GETTING ENOUGH IRON

WITH MY CURRENT TREATMENT OPTION?

IF YOU'RE TAKING IRON SUPPLEMENTS
Even in healthy people, **less than 10% of the iron in iron supplements is absorbed**.*

IF YOU'RE ON AN IV IRON TREATMENT
With IV iron, an infusion **delivers 100% of iron** directly into the bloodstream through a vein.

Even if you've had IV iron infusions, your iron levels may not be where they need to be. Talk to your healthcare professional about how to replenish your iron levels and which IV iron treatment may help.

*Oral iron is typically taken in 300 mg or 320 mg tablets 3-4 times a day. The body is unable to absorb that much iron in the digestive tract at one time, so iron repletion may be possible in smaller oral iron doses over time.

SELECTED SAFETY INFORMATION

What should I tell my healthcare provider before receiving Injectafer® (ferric carboxymaltose)? (cont'd)

Before you receive Injectafer, tell your healthcare provider about all of your medical conditions, including if you:

- Are pregnant or plan to become pregnant. Injectafer may harm your unborn baby. Tell your healthcare provider right away if you become pregnant or think you are pregnant during treatment with Injectafer.

INJECTAFER GETS YOU MORE IRON IN LESS TIME VS ORAL IRON

AND THE MOST IRON OF ANY IV IRON PER COURSE OF TREATMENT*

For patients with iron deficiency anemia (IDA) who can't tolerate iron supplements or who aren't seeing improvements despite being on oral supplements, Injectafer is the only IV iron that provides up to **1500 mg of iron** in 1 course of treatment that happens **over 2 appointments, separated by at least 7 days.**

Injectafer is well studied and widely used, studied in clinical trials with more than **8800 people** and used to treat more than **3 million people** in the United States alone.

OVER
3 MILLION
PEOPLE TREATED IN THE U.S.

ASK ABOUT INJECTAFER

Ask your healthcare professional about Injectafer if you're not getting to goal on your current iron treatment.

*For IDA, one course of treatment is 2 doses of 750 mg separated by at least 7 days. For patients weighing less than 50 kg (110 lb), the recommended dosage is Injectafer 15 mg/kg body weight intravenously in 2 doses separated by at least 7 days per course.

SELECTED SAFETY INFORMATION

What should I tell my healthcare provider before receiving Injectafer? (cont'd)

Before you receive Injectafer, tell your healthcare provider about all of your medical conditions, including if you:

- Are breastfeeding or plan to breastfeed. Injectafer passes into your breast milk. It is not known if Injectafer will harm your baby. Talk to your healthcare provider about the best way to feed your baby during treatment with Injectafer

GETTING YOUR INJECTAFER INFUSIONS

Your healthcare professional has prescribed Injectafer for you because it may be an effective way to replace the iron you need.

WHERE INJECTAFER IS GIVEN

Injectafer is usually given at an infusion center, and administered by a healthcare professional.

CLICK HERE TO VISIT INJECTAFER.COM/LOCATOR

SELECTED SAFETY INFORMATION

What should I tell my healthcare provider before receiving Injectafer? (cont'd)

Tell your healthcare provider about all the medications you take, including prescription and over-the-counter medicines, vitamins, and herbal supplements.

Please click here for Full Prescribing Information and see Important Safety Information on pages 18-19.

injectafer®
ferric carboxymaltose injection

HERE'S WHAT YOU SHOULD KNOW

BEFORE, DURING, AND AFTER HELPING YOU GET REPLENISHED

BEFORE YOUR FIRST INJECTAFER INFUSION

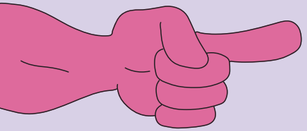
- Call ahead to confirm that the infusion center can provide Injectafer IV iron
- Dress comfortably and eat as you normally would. There are no special dietary requirements
- Bring insurance card, photo identification, and any other information your healthcare professional has asked you to bring

DURING YOUR INFUSION

- Each Injectafer infusion may take about **15 minutes**
- The recommended dosage is **1500 mg**, administered in **2 doses of 750 mg** separated by at least 7 days. Your dosage may vary based on your weight*

AFTER EACH INFUSION

- Afterward, your healthcare provider will monitor you for about 30 minutes for signs of allergic reaction
- In some patients, an increase in blood pressure with dizziness, nausea, or flushing of the face may occur right after a dose of Injectafer. It usually goes away within 30 minutes. Tell your healthcare professional right away if symptoms persist or worsen
- Your iron stores are replenished over time. So, it's important for you to follow up with your healthcare professional to retest your levels and see how Injectafer is working for you



CLICK HERE TO VISIT INJECTAFER.COM/LOCATOR

*For patients weighing less than 50 kg, the recommended dosage is Injectafer 15 mg/kg body weight intravenously in 2 doses separated by at least 7 days per course of treatment.

WHAT SIDE EFFECTS COULD OCCUR IN THE DAYS FOLLOWING AN INFUSION?

- Some patients experience nausea, high blood pressure, flushing, low levels of phosphorous in their blood, dizziness, vomiting, headache, an increase in certain liver enzymes, and pain or bruising at the injection site during or immediately after an infusion
- These are not all of the possible side effects of Injectafer. **Call your healthcare professional for medical advice about side effects**

SEEK MEDICAL ATTENTION IF MORE SERIOUS SIDE EFFECTS ARISE

Serious side effects may include but are not limited to: allergic reactions including itching, hives, wheezing, low blood pressure, and high blood pressure, sometimes with facial flushing, dizziness, or nausea.

SELECTED SAFETY INFORMATION

How will I receive Injectafer?

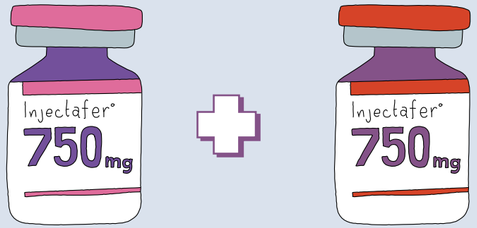
Injectafer is given into your vein (intravenously) by your healthcare provider in 2 doses at least 7 days apart. For certain patients with heart failure, 2 doses may need to be given 6 weeks apart. If your healthcare provider decides it is right for you, Injectafer may be given intravenously by your healthcare provider as a single-dose treatment. Injectafer treatment may be repeated if your healthcare provider decides it is needed.

Please click here for Full Prescribing Information and see Important Safety Information on pages 18-19.

injectafer® ferric carboxymaltose injection

BE SURE TO COMPLETE YOUR INJECTAFER COURSE OF TREATMENT

REMEMBER, INJECTAFER IS GIVEN IN 2 DOSES SEPARATED BY AT LEAST 7 DAYS



- Come back for your **second dose** after at least 7 days
- Remember to schedule or confirm your **next appointment**
- **Track your treatments** and progress using the tables on page 8 of this brochure

SELECTED SAFETY INFORMATION

What are the possible side effects of Injectafer?

Injectafer may cause serious side effects, including:

- **Allergic reactions**. Serious life-threatening allergic reactions that can lead to death have happened in people who receive Injectafer and may include the following signs or symptoms: low blood pressure, feeling dizzy or lightheaded, loss of consciousness, trouble breathing, swelling, fast heartbeat, cold or clammy skin, feet or hands turn blue, itching, rash, hives, and/or wheezing. Your healthcare provider will watch you during and for at least 30 minutes after you receive Injectafer. Tell your healthcare provider right away if you develop any signs or symptoms of allergic reactions during or after treatment with Injectafer

REMEMBER THE 3 "C"s

CHECK

your hemoglobin (Hb) blood test results on your iron panel

COMPARE

vs healthy levels with your healthcare professional's input

CONSULT

with your healthcare professional about switching to Injectafer after oral iron failure if your Hb level is not at a healthy level*

CLICK HERE TO VISIT INJECTAFER.COM/INFO

*Normal lab values may vary based on patient characteristics/ comorbidities and by laboratory.

Please click here for Full Prescribing Information and see Important Safety Information on pages 18-19.

injectafer® ferric carboxymaltose injection

TWO WAYS TO HELP YOU PAY FOR INJECTAFER TREATMENTS

If you have an Injectafer prescription, you may be able to get help with your out-of-pocket costs.

1

IF INJECTAFER IS COVERED BY YOUR COMMERCIAL INSURANCE, BUT YOU HAVE A CO-PAY

INJECTAFER SAVINGS PROGRAM*

The Injectafer Savings Program may help eligible patients with their Injectafer prescription out-of-pocket responsibility.

PATIENTS RECEIVE EACH DOSE FOR AS LITTLE AS $50

For eligible patients:

- Assistance of up to $500 per dose
- Enrollment is valid for 2 courses of treatment per 12-month period

HOW TO ENROLL

Ask your healthcare professional to enroll you prior to receiving Injectafer treatment. If your healthcare professional cannot enroll you, you can enroll yourself in one of two ways:

- The best way to enroll is by visiting **injectafercopay.com**

OR

- **Call Daiichi Sankyo Access Central (1-866-437-4669)**

2

IF YOU DO NOT HAVE INSURANCE TO COVER TREATMENT OR ARE COMMERCIALLY UNDERINSURED

PATIENT ASSISTANCE PROGRAM†

The Patient Assistance Program was created to help patients who lack health insurance or are commercially underinsured and cannot afford therapy.

DAIICHI SANKYO ACCESS CENTRAL

1-866-4-DSI-NOW (1-866-437-4669) is available.

*The Injectafer Savings Program is only available for patients aged 1 year or older who are commercially insured. Insurance out-of-pocket must be over $50 per dose. Additional restrictions may apply. Please see full Terms and Conditions at DSIAccessCentral.com/patient/home.

†Terms and conditions apply.

Please click here for Full Prescribing Information and see Important Safety Information on pages 18-19.

injectafer® ferric carboxymaltose injection

Injectafer® (ferric carboxymaltose injection) is available by prescription only. Ask your healthcare provider if Injectafer is right for you.

What is Injectafer?

Injectafer is a prescription iron replacement medicine administered only by or under the supervision of your healthcare provider. Injectafer is injected into your vein to treat iron deficiency anemia in adults and children 1 year of age and older. Injectafer should be used only if you have not responded well to treatment with oral iron, or if you are intolerant to oral iron treatment. It is also used to treat iron deficiency anemia in adults with chronic kidney disease who are not receiving dialysis. Injectafer is used to improve the ability to exercise (exercise capacity) in adult patients with iron deficiency and mild to moderate heart failure. It is not known if Injectafer is safe and effective in children with iron deficiency anemia who are under 1 year of age or in children with iron deficiency and mild to moderate heart failure to improve exercise capacity.

IMPORTANT SAFETY INFORMATION

Who should not receive Injectafer?

You should not receive Injectafer if you are allergic to ferric carboxymaltose or any of the other ingredients in Injectafer. The active ingredient in Injectafer is ferric carboxymaltose, the inactive ingredients are: water for injection, sodium hydroxide or hydrochloric acid.

What should I tell my healthcare provider before receiving Injectafer?

Before you receive Injectafer, tell your healthcare provider about all of your medical conditions, including if you:

- Have had an allergic reaction to iron given into your vein
- Have a history of trouble absorbing certain vitamins or phosphate in your body
- Have inflammatory bowel disease
- Have hyperparathyroidism
- Have low vitamin D levels
- Have high blood pressure
- Have previously received Injectafer
- Are pregnant or plan to become pregnant. Injectafer may harm your unborn baby. Tell your healthcare provider right away if you become pregnant or think you are pregnant during treatment with Injectafer.
- Are breastfeeding or plan to breastfeed. Injectafer passes into your breast milk. It is not known if Injectafer will harm your baby. Talk to your healthcare provider about the best way to feed your baby during treatment with Injectafer

Tell your healthcare provider about all the medications you take, including prescription and over-the-counter medicines, vitamins, and herbal supplements.

How will I receive Injectafer?

Injectafer is given into your vein (intravenously) by your healthcare provider in 2 doses at least 7 days apart. For certain patients with heart failure, 2 doses may need to be given 6 weeks apart. If your healthcare provider decides it is right for you, Injectafer may be given intravenously by your healthcare provider as a single-dose treatment. Injectafer treatment may be repeated if your healthcare provider decides it is needed.

Please click here for Full Prescribing Information for Injectafer.

What are the possible side effects of Injectafer?

Injectafer may cause serious side effects, including:

- **Allergic reactions**. Serious life-threatening allergic reactions that can lead to death have happened in people who receive Injectafer and may include the following signs or symptoms: low blood pressure, feeling dizzy or lightheaded, loss of consciousness, trouble breathing, swelling, fast heartbeat, cold or clammy skin, feet or hands turn blue, itching, rash, hives, and/or wheezing. Your healthcare provider will watch you during and for at least 30 minutes after you receive Injectafer. Tell your healthcare provider right away if you develop any signs or symptoms of allergic reactions during or after treatment with Injectafer
- **Symptoms of low blood phosphate levels**. Injectafer may cause low levels of phosphate in your blood that may be serious and can lead to softening of your bones and broken bones (fractures), especially in people who have received multiple Injectafer treatments. Your healthcare provider may check your blood phosphate levels before a repeat treatment with Injectafer if you are at risk for low blood phosphate levels. If a repeat treatment is needed within 3 months of your last treatment, your healthcare provider should check your blood phosphate levels. Tell your healthcare provider if you develop any of the following signs or symptoms of low blood phosphate levels during treatment with Injectafer: feeling very tired, muscle weakness or pain, bone or joint pain, broken bones
- **High blood pressure**. High blood pressure, sometimes with redness and warmth of the face (facial flushing), dizziness, or nausea, has happened during treatment with Injectafer. Your healthcare provider will check your blood pressure and check for any signs and symptoms of high blood pressure after you receive Injectafer

The most common side effects of Injectafer include:

- In adults: nausea, high blood pressure, flushing, injection site reactions, skin redness, low levels of phosphate in your blood, and dizziness.
- In children: low levels of phosphate in your blood, injection site reactions, rash, headache, and vomiting

These are not all of the possible side effects of Injectafer.

Call your healthcare provider for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

General information about the safe and effective use of Injectafer

Medicines are sometimes prescribed for purposes other than those listed in a Patient Information leaflet. You can ask your pharmacist or healthcare provider for information about Injectafer that is written for health professionals.

To report side effects, contact American Regent at 1-800-734-9236 or E-mail: pv@americanregent.com or Fax: 1-610-650-0170.

You may also report side effects to the FDA at 1-800-332-1088 or www.fda.gov/medwatch.

The risk information provided here is not comprehensive. To learn more about Injectafer, talk with your healthcare provider or pharmacist. The FDA-approved product labeling can be found at www.injectafer.com/pdf/pi.pdf or call 1-800-645-1706.

injectafer®
ferric carboxymaltose injection

Choose

INJECTAFER, AN IV IRON TREATMENT

- Provides **more iron in less time** vs oral iron and the **most iron** of any IV iron per course of treatment for IDA*
- **Delivers 100% of iron** directly into the bloodstream
- Used to treat more than **3 million people** in the United States alone

CLICK HERE TO VISIT INJECTAFER.COM/INFO

*For iron deficiency anemia (IDA), one course of treatment is 2 doses of 750 mg separated by at least 7 days. For patients weighing less than 50 kg (110 lb), the recommended dosage is Injectafer 15 mg/kg body weight intravenously in 2 doses separated by at least 7 days per course.

SELECTED SAFETY INFORMATION ABOUT INJECTAFER

Who should not receive Injectafer? You should not receive Injectafer if you are allergic to ferric carboxymaltose or any of the other ingredients in Injectafer. The active ingredient in Injectafer is ferric carboxymaltose, the inactive ingredients are: water for injection, sodium hydroxide or hydrochloric acid.

Please click here for Full Prescribing Information and see Important Safety Information on pages 18-19.

Daiichi-Sankyo

To contact us with questions or concerns about a Daiichi Sankyo product, please call us: 1-877-4DS-PROD (1-877-437-7763).

American Regent, Inc. is a member of the Daiichi Sankyo Group. Injectafer® and the Injectafer® logo are trademarks of Vifor (International) Inc., Switzerland. Injectafer® is manufactured under license from Vifor (International) Inc., Switzerland. Trademarks not owned by American Regent, Inc. or Vifor (International) are the property of their respective owners.

 PP-US-IN-4702 03/25

injectafer® ferric carboxymaltose injection